LISTING OF THE CLAIMS

Docket No.: IFM-005CP4CN2RCE

This listing of claims will replace all prior versions of the claims and listing of the claims in the application:

1.-13. (Cancelled)

14. **(Previously Presented)** A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:

$$R_4 - R_3 - CH - N$$
 $R_2 - CH_2 - C \equiv CH$
(II)

in which

 R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

R₂ is hydrogen or alkyl;

R₃ is a bond or methylene; and

R₄ is aryl or aralkyl; or

 R_2 and R_4 - R_3 are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group;

and pharmaceutically acceptable salts thereof.

15. **(Previously Presented)** A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:

$$R_4 - R_3 - CH - N$$
 $R_2 R_5 - C \equiv CH$
(III)

in which

 R_2 is hydrogen or alkyl;

R₃ is a bond or methylene; and

R₄ is aryl or aralkyl; or

 R_2 and R_4 - R_3 are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group; and

R₅ is alkylene, alkenylene, alkynylene and alkoxylene; and pharmaceutically acceptable salts thereof.

16. (**Previously Presented**) A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:

$$\stackrel{A_n}{\longleftarrow} -CH_2 - CH - N$$
 $\stackrel{CH_2}{\longleftarrow} -CH_2 - C \equiv CH$
 $\stackrel{CH_3}{\longleftarrow} -CH_2 - C \equiv CH$
 $\stackrel{(IV)}{\longleftarrow} -CH_2 - C \equiv CH$

in which

 R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

A is a substituent independently selected for each occurrence from the group consisting of halogen, hydroxyl, alkyl, alkoxyl, cyano, nitro, amino, carboxyl, -CF₃, or azido;

n is 0 or an integer from 1 to 5;

and pharmaceutically acceptable salts thereof.

17.-20. (Cancelled)

- 21. **(Previously Presented)** The method of claim 14, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
- 22. **(Previously Presented)** The method of claim 21, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
- 23. **(Previously Presented)** The method of claim 22, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.

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24. **(Previously Presented)** The method of claim 21, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.

- 25. (Previously Presented) The method of claim 14, wherein said subject is a human.
- 26. **(Previously Presented)** The method of claim 15, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
- 27. **(Previously Presented)** The method of claim 26, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
- 28. **(Previously Presented)** The method of claim 26, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.
- 29. **(Previously Presented)** The method of claim 28, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
- 30. **(Previously Presented)** The method of claim 15, wherein said subject is a human.
- 31. **(Previously Presented)** The method of claim 16, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
- 32. **(Previously Presented)** The method of claim 31, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
- 33. **(Previously Presented)** The method of claim 31, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.

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34. **(Previously Presented)** The method of claim 31, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.

35. (**Previously Presented**) The method of claim 16, wherein said subject is a human.